Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Currently Amended) A pharmaceutical composition ~~for moulded capsules~~ comprising Eudragit 4135F present in an amount of about 20 to 90% w/w; a lubricant present in an amount of 0 to about 30% w/w; a dissolution modifying excipient present in an amount of about 2.5 to about 70% w/w, and optionally a surfactant present in an amount of 0 to 10%, a plasticizer present in an amount of 0 to 10% w/w and/or a processing agent present in an amount of 0 to about 10% w/w for moulded capsule, shell or linker components.

2. (Original) The composition according to Claim 1 wherein the Eudragit 4135F is present in an amount of about 50 to 90% w/w.

3. (Original) The composition according to Claim 1 which comprises a surfactant which is present in an amount of less than 5% w/w.

4. (Original) The composition according to Claim 3 wherein the surfactant is sodium dodecyl sulphate or is a block copolymer of ethylene oxide and propylene oxide.

5. (Original) The composition according to Claim 4 wherein the surfactant is sodium dodecyl sulphate is present in an amount of less than 2% w/w.

6. (Original) The composition according to Claim 4 wherein the surfactant is a block copolymer of ethylene oxide and propylene oxide.

7. (Original) The composition according to Claim 1 wherein the lubricant is present in an amount of about 10 to 30% w/w.

8. (Currently amended) The composition according to Claim 1 wherein the lubricant is stearyl alcohol, glycerol monostearate ~~monosterate~~ (GMS), talc, magnesium stearate, silicon dioxide, amorphous silicic acid, or fumed silica; and combinations or mixtures thereof.

9. (Original) The composition according to Claim 8 wherein the lubricant is stearyl alcohol.

10. (Original) The composition according to Claim 9 wherein the stearyl alcohol is present from about 10 to about 15% w/w.

11. (Original) The composition according to Claim 1 wherein the dissolution modifying excipient is a swellable solid which is ethyl cellulose, cellulose acetate phthalate; hydroxypropyl cellulose, hydroxypropylmethyl cellulose, hydroxypropylmethyl cellulose phthalate, or other hydroxyalkylcellulose derivative; and combinations or mixtures thereof.

12. (Original) The composition according to Claim 11 wherein the dissolution modifying excipient is hydroxypropylmethyl cellulose, hydroxypropylmethyl cellulose phthalate, or hydroxypropyl cellulose.

13. (Currently amended) The composition according to Claim 12 wherein the swellable solid is present in an amount of about 10 to 50% w/w.

14. (Currently amended) The composition according to Claim 1 wherein the dissolution modifying excipient is xylitol, mannitol, lactose, pregelatinized starch ~~Starch 1500~~, sodium chloride, sodium starch glycollate, croscarmellose sodium, crospovidone (cross-linked polyvinyl pyrrolidone), copovidone , polyvinyl pyrrolidone, ~~copovidone~~; and combinations or mixtures thereof.

15. (Currently amended) The composition according to Claim 14 wherein the dissolution modifying excipient is present in an amount of about 40 to 70% w/w.

16. (Currently amended) The composition according to Claim 11 wherein the dissolution modifying excipient is a combination of a swellable solid and lactose, sodium starch glycollate, croscarmellose sodium, copovidone, crospovidone (cross-linked polyvinyl pyrrolidone), or polyvinyl pyrrolidone ~~copovidone~~.

17. (Original) The composition according to Claim 16 wherein the dissolution modifying excipient is hydroxypropylcellulose and lactose.

18. (Original) The composition according to Claim 1 wherein the surfactant is a block copolymer of ethylene oxide and propylene oxide, lecithin, sodium dioctyl

sulfosuccinate, sodium dodecyl sulphate, Polyoxyl 40 hydrogenated castor oil, polyoxyethylene sorbitan fatty acid esters, the sorbitan fatty acid esters, polyethylene glycol, Vitamin E-TPGS® (d-alpha-tocopheryl polyethylene glycol 1000 succinate), sucrose fatty acid ester; and combinations and mixtures thereof.

19. (Currently Amended) The composition according Claim 18 wherein the dissolution modifying excipient is a combination of a swellable solid and lactose, sodium starch glycollate, croscarmellose sodium, copovidone, crospovidone (cross-linked polyvinyl pyrrolidone), or polyvinyl pyrrolidone ~~copovidone~~.

20. (Original) The composition according to Claim 1 wherein the plasticizer is triethyl citrate (TEC), tributyl citrate, acetyl triethyl citrate (ATEC), acetyl tributyl citrate (ATBC), dibutyl phthalate, dibutyl sebacate (DBS), diethyl phthalate, vinyl pyrrolidone glycol triacetate, polyethylene glycol, polyoxyethylene sorbitan monolaurate, propylene glycol, or castor oil; and combinations or mixtures thereof.

21. (Original) The composition according to Claim 1 wherein the processing agent is talc.

22. (Original) The composition according to Claim 21 wherein the processing agent is present in an amount of about 1 to about 5 % w/w.

23. (Original) The composition according to Claim 1 which further comprises an absorption enhancer.

24. (Original) The composition according to Claim 23 wherein the absorption enhancer is chitosan, lecithin, lectin, a sucrose fatty acid ester, Vitamin E-TPGS; and combinations or mixtures thereof.

25. (Original) The composition according to Claim 1 wherein the Eudragit 4135F is present in an amount of about 50 to 90% w/w, the lubricant is stearyl alcohol, and the dissolution modifying excipient is hydroxypropylmethylcellulose, hydroxypropylcellulose, or a hydroxylalkyl cellulose derivative or salt thereof.

26. (Original) The composition according to Claim 25 wherein the dissolution modifying excipient also includes a disintegrant.

27. (Currently Amended) The composition according to Claim 26 wherein the disintegrant is sodium starch glycollate, croscarmellose sodium, copovidone, crospovidone (cross-linked polyvinyl pyrrolidone), or polyvinyl pyrrolidone ~~copovidone~~, or a combination or mixture thereof.

28. (Original) The composition according to Claim 25 wherein the dissolution modifying excipient also includes a wicking agent.

29. (Original) The composition according to Claim 28 wherein the wicking agent is lactose.

30. (Original) The composition according to Claim 25 wherein the processing aid is talc.

31. (Currently Amended) The pharmaceutical composition according to Claim 1 which is:

Example #	Formulation ~~(% w/w)~~	% w/w
1.	Eudragit 4135F	75.0
	Stearyl alcohol	5.0
	Croscarmellose sodium	20.0
2.	Eudragit 4135F	75.0
	Stearyl alcohol	5.0
	Sodium starch glycollate	20.0
3.	Eudragit 4135F	85.0
	Stearyl alcohol	5.0
	Xylitol	10.0
4.	Eudragit 4135F	75.0
	Stearyl alcohol	5.0
	Croscarmellose sodium	10.0
	Xylitol	10.0
5.	Eudragit 4135F	75.0
	Stearyl alcohol	5.0
	Mannitol	10.0
	Sodium starch glycollate	10.0
6.	Eudragit 4135F	65.0
	Stearyl alcohol	5.0
	Mannitol	10.0
	Sodium starch glycollate	20.0

7.	Eudragit 4135F Stearyl alcohol Sodium starch glycollate Lactose monohydrate	80.0 5.0 10.0 5.0
8.	Eudragit 4135F Stearyl alcohol Sodium starch glycollate Lactose monohydrate	75.0 5.0 10.0 10.0
9.	Eudragit 4135F Stearyl alcohol Lactose monohydrate	85.0 5.0 10.0
10.	Eudragit 4135F Stearyl alcohol Lactose monohydrate	75.0 5.0 20.0
11.	Eudragit 4135F Stearyl alcohol Sodium starch glycollate Lactose monohydrate	80.0 5.0 5.0 10.0
12.	Eudragit 4135F Stearyl alcohol Sodium starch glycollate Lactose monohydrate	70.0 5.0 5.0 20.0
13.	Eudragit 4135F Stearyl alcohol Mannitol Sodium starch glycollate	75.0 10.0 7.5 7.5
14.	Eudragit 4135F Stearyl alcohol ~~Starch 1500~~ Pregelatinized Starch	80.0 5.0 10.0
15.	Eudragit 4135F Stearyl alcohol ~~Starch 1500~~ Pregelatinized Starch	85.0 5.0 15.0
16.	Eudragit 4135F Stearyl alcohol ~~Starch 1500~~ Pregelatinized Starch Lactose monohydrate	80.0 5.0 10.0 5.0
17.	Eudragit 4135F Stearyl alcohol ~~Kollidon CL~~ Cross linked polyvinyl pyrrolidone	85.0 5.0 10.0
18.	Eudragit 4135F Stearyl alcohol Sodium starch glycollate Lactose monohydrate	80.0 5.0 10.0 5.0

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19.	Eudragit 4135F Stearyl alcohol Sodium starch glycollate Lactose monohydrate	75.0 10.0 10.0 5.0
20.	Eudragit 4135F Stearyl alcohol Sodium chloride Lactose monohydrate	85.0 5.0 5.0 5.0
21.	Eudragit 4135F Stearyl alcohol ~~Klucel LF~~ Hydroxypropyl cellulose Lactose monohydrate	85.0 5.0 5.0 5.0
22.	Eudragit 4135F Stearyl alcohol Hydroxypropylmethyl cellulose Lactose monohydrate	85.0 5.0 5.0 5.0
23.	Eudragit 4135F Stearyl alcohol Hydroxypropylmethyl cellulose Lactose monohydrate	80.0 10.0 5.0 5.0
24.	Eudragit 4135F Stearyl alcohol Sodium starch glycollate Lactose monohydrate	80.0 10.0 5.0 5.0
25.	Eudragit 4135F Stearyl alcohol Hypromellose phthallate Lactose monohydrate	80.0 10.0 5.0 5.0
26.	Eudragit 4135F Stearyl alcohol Low substituted hydroxypropyl cellulose Lactose monohydrate	80.0 10.0 5.0 5.0
27.	Eudragit 4135F Stearyl alcohol Hydroxypropylmethyl cellulose	90.0 5.0 5.0
28.	Eudragit 4135F Stearyl alcohol Lactose monohydrate	90.0 5.0 5.0
29.	Eudragit 4135F Stearyl alcohol Hydroxypropylmethyl cellulose Lactose monohydrate	73.0 12.0 10.0 5.0
30.	Eudragit 4135F Sodium dodecyl sulphate Croscarmellose sodium	84.0 1.0 15%

31.	Eudragit 4135F Sodium dodecyl sulphate Croscarmellose sodium Sodium starch glycollate	79.0 1.0 10% 10%
32.	Eudragit 4135F Croscarmellose sodium Sodium starch glycollate	80.0 10% 10%
33.	Eudragit 4135F Sodium dodecyl sulphate Croscarmellose sodium Sodium starch glycollate	69.0 1.0 15% 15%
34.	Eudragit 4135F ~~Pluronic F68~~ Polyoxypropylene-polyoxyethylene block copolymer Sodium starch glycollate	79.0 1.0 20%
35.	Eudragit 4135F ~~Pluronic F127~~ Polyoxypropylene-polyoxyethylene block copolymer Sodium starch glycollate	79.0 1.0 20%

32. (Currently Amended) A pharmaceutical composition for molded capsule shells comprising:

Components	# (1) w/w	(2) w/w	(3) w/w	(4) w/w	(5) w/w	(6) w/w	(7) w/w
Eudragit 4135F	45%	35%	25%	15%	75%	65%	55%
Stearyl Alcohol	10%	10%	10%	10%	10%	10%	10%
Lactose	5%	5%	5%	5%	5%	5%	5%
~~Klucel LF~~ Hydroxypropyl Cellulose	40%	50%	60%	70%	10%	20%	30%
Total	100%	100%	100%	100%	100%	100%	100%

33. (Currently Amended) A pharmaceutical composition for molded capsule shells comprising:

Components	# (1) w/w	(2) w/w	(3) w/w	(4) w/w	(5) w/w	(6) w/w
Eudragit 4135F	63%	62.9%	62.75%	52%	42%	62%
Croscarmellose sodium	10%	10%	10%	15%	20%	5%
Sodium starch glycollate	10%	10%	10%	15%	20%	5%
Stearyl alcohol	12%	12%	12%	12%	12%	12%
Hydroxypropyl-methylcellulose	5%	5%	5%	5%	5%	15%
Sodium Dodecyl Sulphate	0%	0.1%	0.25%	1%	1%	1%.

34. (Currently Amended) A pharmaceutical composition for molded capsule shells comprising:

Components	#(1) w/w	(2) w/w	(3) w/w	(4) w/w	(5) w/w	(6) w/w	(7) w/w
Eudragit 4135F	45%	35%	25%	15%	75%	65%	55%
Stearyl Alcohol	10%	10%	10%	10%	10%	10%	10%
Lactose	5%	5%	5%	5%	5%	5%	5%
~~Klucel LF~~ Hydroxypropyl cellulose	40%	50%	60%	70%	10%	20%	30%
Total	100%	100%	100%	100%	100%	100%	100%

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35. (Currently Amended) A pharmaceutical composition for molded capsule shells comprising:

Example #	Formulation	%w/w
1.	Eudragit 4135F	73.0
	Hydroxypropylmethyl cellulose	10.0
	Lactose (regular)	5.0
	Glyceryl monostearate	12.0
2.	Eudragit 4135F	53.0
	Hydroxypropylmethyl cellulose	10.0
	Lactose (regular)	5.0
	Hydroxypropylmethyl cellulose phthallate	20.0
	Stearyl alcohol	12.0
3.	Eudragit 4135F	20.0
	Hydroxypropylmethyl cellulose	10.0
	Hydroxypropylmethyl cellulose phthallate	20.0
	Stearyl alcohol	12.0
4.	Eudragit 4135F	68.0
	Hydroxypropylmethyl cellulose	10.0
	Lactose (regular)	5.0
	Sodium dodecyl sulphate	5.0
	Stearyl alcohol	12.0
5.	Eudragit 4135F	72.0
	Hydroxypropylmethyl cellulose	10.0
	Lactose (regular)	5.0
	Sodium dodecyl sulphate	1.0
	Stearyl alcohol	12.0
6.	Eudragit 4135F	71.0
	Hydroxypropylmethyl cellulose	10.0
	Lactose (regular)	5.0
	Sodium dodecyl sulphate	2.0
	Stearyl alcohol	12.0
7.	Eudragit 4135F	62.0
	Sodium starch glycollate	20.0
	Lactose (regular)	5.0
	Sodium dodecyl sulphate	1.0
	Stearyl alcohol	12.0
8.	Eudragit 4135F	62.0
	Sodium starch glycollate	20.0
	Lactose (regular)	5.0
	Sodium dodecyl sulphate	1.0
	Stearyl alcohol	12.0
9.	Eudragit 4135F	72.0
	Sodium starch glycollate	10.0
	Lactose (regular)	5.0
	Sodium dodecyl sulphate	1.0
	Stearyl alcohol	12.0

10.	Eudragit 4135F Croscarmellose sodium Lactose (regular) Sodium dodecyl sulphate Stearyl alcohol	62.0 20.0 5.0 1.0 12.0
11.	Eudragit 4135F Sodium starch glycollate Hydroxypropylmethyl cellulose Sodium dodecyl sulphate Stearyl alcohol	62.0 20.0 5.0 1.0 12.0
12.	Eudragit 4135F Hydroxypropylmethyl cellulose phthallate Hydroxypropylmethyl cellulose Sodium dodecyl sulphate Stearyl alcohol	62.0 20.0 5.0 1.0 12.0
13.	Eudragit 4135F Sodium starch glycollate Hydroxypropylmethyl cellulose Sodium dodecyl sulphate Stearyl alcohol	62.5 20.0 5.0 0.5 12.0
14.	Eudragit 4135F Croscarmellose sodium Sodium starch glycollate Hydroxypropylmethyl cellulose Sodium dodecyl sulphate Stearyl alcohol	62.0 10.0 10.0 5.0 1.0 12.0
15.	Eudragit 4135F Croscarmellose sodium Hydroxypropylmethyl cellulose Sodium dodecyl sulphate Stearyl alcohol	67.0 15.0 5.0 1.0 12.0
16.	Eudragit 4135F Croscarmellose sodium Hydroxypropylmethyl cellulose Sodium dodecyl sulphate Stearyl alcohol	72.0 10.0 5.0 1.0 12.0
17.	Eudragit 4135F Croscarmellose sodium Hydroxypropylmethyl cellulose Sodium dodecyl sulphate Stearyl alcohol	77.0 5.0 5.0 1.0 12.0
18.	Eudragit 4135F Croscarmellose sodium Sodium starch glycollate Hydroxypropylmethyl cellulose Sodium dodecyl sulphate Stearyl alcohol	52.0 15.0 15.0 5.0 1.0 12.0

19.	Eudragit 4135F Croscarmellose sodium Sodium starch glycollate Hydroxypropylmethyl cellulose Sodium dodecyl sulphate Stearyl alcohol	42.0 20.0 20.0 5.0 1.0 12.0
20.	Eudragit 4135F Croscarmellose sodium Sodium starch glycollate Hydroxypropylmethyl cellulose Sodium dodecyl sulphate Stearyl alcohol	42.0 20.0 20.0 5.0 1.0 12.0
21.	Eudragit 4135F Croscarmellose sodium Sodium starch glycollate Hydroxypropylmethyl cellulose Sodium dodecyl sulphate Stearyl alcohol	62.0 5.0 5.0 15.0 1.0 12.0
22.	Eudragit 4135F Croscarmellose sodium Sodium starch glycollate Hydroxypropylmethyl cellulose Sodium dodecyl sulphate Stearyl alcohol	62.9 10.0 10.0 5.0 0.1 12.0

36. (Cancelled)

37. (Cancelled)

38. (Currently Amended) The composition according to Claim 1 wherein the lubricant is stearyl alcohol present in an amount of 10 to 15% w/w, the surfactant ~~lubricant~~ is SDS or a ~~present in an amount of a~~ block copolymer of ethylene oxide and propylene oxide present in an amount ~~at~~ less than 5% w/w; a dissolution modifying excipient selected from ~~is~~ HPC, HPMC, sodium starch glycollate, croscarmellose sodium, copovidone, or lactose, and combinations or mixtures thereof, present in an amount of about 2.5 to about 70% w/w.

39. (Currently Amended) A composition according to Claim 1 that is in the form of an ~~An~~ injection molded capsule shell, linker or spacer ~~having a composition according to Claim 1~~.

40. (Currently Amended) A composition according to Claim 1 that is in the form of a ~~A~~ multicomponent injection molded capsule shell, linker or spacer ~~having a composition according to Claim 1~~.

41. (Currently Amended) A composition according to Claim 1 that is in the form of a ~~A~~ welded multicomponent injection molded capsule shell, linker or spacer ~~having a composition according to Claim 1~~.

42 to 70 (cancelled).

71. (new) The composition according to Claim 1 wherein the

	Dissolution Modifier	Lubricant	Surfactant
1.	Hydroxypropylmethylcellulose (5%w/w)	Stearyl alcohol (12%w/w)	None; or
2.	Hydroxypropylmethylcellulose (10%w/w), and HPMCphthalate (20%w/w)	Stearyl alcohol (12%w/w)	None; or
3.	Hydroxypropylmethylcellulose (10%), and Lactose (5%)	Stearyl alcohol (12%)	None; or
4.	Hydroxypropylmethylcellulose (5%)	Stearyl alcohol (12%)	SDS (1%) or Sodium Starch Glycollate (20%) or Tween or a polyoxypropylene-polyoxyethylene block copolymer.

72. (New) The composition according to Claim 1 which is

Example #	Formulation	% w/w
1.	Eudragit 4135F	77.0
	Sodium Dodecyl Sulphate	1.0
	Croscarmellose sodium	5.0
	Stearyl Alcohol	12.0
	Hydroxypropylmethyl Cellulose	5.0
2.	Eudragit 4135F	68.0
	Croscarmellose sodium	15.0
	Stearyl Alcohol	12.0
	Hydroxypropylmethyl Cellulose	5.0

3.	Eudragit 4135F	62.0
	Sodium Dodecyl Sulphate	1.0
	Croscarmellose sodium	10.0
	Sodium Starch Glycollate	10.0
	Stearyl Alcohol	12.0
	Hydroxypropylmethyl Cellulose	5.0
4.	Eudragit 4135F	63.0
	Croscarmellose sodium	10.0
	Sodium Starch Glycollate	10.0
	Stearyl Alcohol	12.0
	Hydroxypropylmethyl Cellulose	5.0
5.	Eudragit 4135F	52.0
	Sodium Dodecyl Sulphate	1.0
	Croscarmellose sodium	15.0
	Sodium Starch Glycollate	15.0
	Stearyl Alcohol	12.0
	Hydroxypropylmethyl Cellulose	5.0
6.	Eudragit 4135F	62.0
	Polyoxypropylene-polyoxyethylene block copolymer	1.0
	Sodium Starch Glycollate	20.0
	Stearyl Alcohol	12.0
	Hydroxypropylmethyl Cellulose	5.0
7.	Eudragit 4135F	62.0
	polyoxypropylene-polyoxyethylene block copolymer	1.0
	Sodium Starch Glycollate	20.0
	Stearyl Alcohol	12.0
	Hydroxypropylmethyl Cellulose	5.0
8.	Eudragit 4135F	62.0
	Stearyl Alcohol	12.0
	Croscarmellose sodium	5.0
	Sodium Starch Glycollate	5.0
	Hydroxypropylmethyl Cellulose	15.0
	Sodium Dodecyl Sulphate	1.0
9.	Eudragit 4135F	42.0
	Stearyl Alcohol	12.0
	Croscarmellose sodium	20.0
	Sodium Starch Glycollate	20.0
	Hydroxypropylmethyl Cellulose	5.0
	Sodium Dodecyl Sulphate	1.0
10.	Eudragit 4135F	47.0
	Stearyl Alcohol	12.0
	Sodium Starch Glycollate	10.0
	Hydroxypropylmethyl Cellulose	30.0
	Sodium Dodecyl Sulphate	1.0

73. (New) A pharmaceutical composition comprising Eudragit 4135F present in an amount of about 20 to 90% w/w; a lubricant present in an amount of 10 to about 30% w/w; a dissolution modifying excipient present in an amount of about 2.5 to about 70% w/w, and optionally a surfactant present in an amount of less than 5% w/w, a plasticizer present in an amount of 0 to 10% w/w and/or a processing agent present in an amount of 0 to about 10% w/w for moulded capsule, shell or linker components.

74. (New) The composition according to Claim 73 wherein the Eudragit 4135F is present in an amount of about 50 to 90% w/w.

75. (New) The composition according to Claim 73 wherein the surfactant is a block copolymer of ethylene oxide and propylene oxide, lecithin, sodium dioctyl sulfosuccinate, sodium dodecyl sulphate, Polyoxyl 40 hydrogenated castor oil, polyoxyethylene sorbitan fatty acid esters, the sorbitan fatty acid esters, polyethylene glycol, Vitamin E-TPGS® (d-alpha-tocopheryl polyethylene glycol 1000 succinate), sucrose fatty acid ester; and combinations and mixtures thereof.

76. (New) The composition according to Claim 75 wherein the surfactant is sodium dodecyl sulphate is present in an amount of less than 2% w/w.

77. (New) The composition according to Claim 75 wherein the surfactant is sodium dodecyl sulphate or is a block copolymer of ethylene oxide and propylene oxide.

78. (New) The composition according to Claim 77 wherein the surfactant is sodium dodecyl sulphate is present in an amount of less than 2% w/w.

79. (New) The composition according to Claim 73 wherein the surfactant is a block copolymer of ethylene oxide and propylene oxide.

80. (New) The composition according to Claim 79 wherein the surfactant is sodium dodecyl sulphate is present in an amount of less than 2% w/w.

81. (New) The composition according to Claim 73 wherein the lubricant is stearyl alcohol, glycerol monostearate (GMS), talc, magnesium stearate, silicon dioxide, amorphous silicic acid, or fumed silica; and combinations or mixtures thereof.

82. (New) The composition according to Claim 81 wherein the lubricant is stearyl alcohol.

83. (New) The composition according to Claim 82 wherein the stearyl alcohol is present from about 10 to about 15% w/w.

84. (New) The composition according to Claim 73 wherein the dissolution modifying excipient is a swellable solid which is ethyl cellulose, cellulose acetate phthalate; hydroxypropyl cellulose, hydroxypropylmethyl cellulose, hydroxypropylmethyl cellulose phthalate, or other hydroxyalkylcellulose derivative; and combinations or mixtures thereof.

85. (New) The composition according to Claim 84 wherein the swellable solid is present in an amount of about 10 to 50% w/w.

86. (New) The composition according to Claim 84 wherein the dissolution modifying excipient is hydroxypropylmethyl cellulose, hydroxypropylmethyl cellulose phthalate, or hydroxypropyl cellulose.

87. (New) The composition according to Claim 86 wherein the swellable solid is present in an amount of 10 to 50% w/w.

88. (New) The composition according to Claim 73 wherein the dissolution modifying excipient is xylitol, mannitol, lactose, pregelatinized starch, sodium chloride, sodium starch glycollate, croscarmellose sodium, crospovidone (cross-linked polyvinyl pyrrolidone), copovidone, polyvinyl pyrrolidone; and combinations or mixtures thereof.

89. (New) The composition according to Claim 88 wherein the dissolution modifying excipient is present in an amount of about 40 to 70% w/w.

90. (New) The composition according to Claim 89 wherein the dissolution modifying excipient is a combination of a swellable solid and lactose, sodium starch glycollate, croscarmellose sodium, copovidone, crospovidone (cross-linked polyvinyl pyrrolidone), or polyvinyl pyrrolidone.

91. (New) The composition according to Claim 90 wherein the dissolution modifying excipient is hydroxypropylcellulose and lactose.

92. (New) The composition according Claim 73 wherein the dissolution modifying excipient is a combination of a swellable solid and lactose, sodium starch glycollate, croscarmellose sodium, copovidone, or crospovidone (cross-linked polyvinyl pyrrolidone).

93. (New) The composition according to Claim 73 wherein the processing agent is talc.

94. (New) The composition according to Claim 93 wherein the processing agent is present in an amount of about 1 to about 5 % w/w.

95. (New) The composition according to Claim 91 wherein the processing agent is talc and is present in an amount of about 1 to about 5 % w/w.

96. (New) The composition according to Claim 73 which further comprises an absorption enhancer.

97. (New) The composition according to Claim 96 wherein the absorption enhancer is chitosan, lecithin, lectin, a sucrose fatty acid ester, Vitamin E-TPGS; and combinations or mixtures thereof.

98. (New) A pharmaceutical composition comprising Eudragit 4135F is present in an amount of about 50 to about 90% w/w, the lubricant is stearyl alcohol, present in an amount of about 10 to about 15% w/w, and the dissolution modifying excipient is hydroxypropylmethylcellulose, hydroxypropylcellulose, or a hydroxylalkyl cellulose derivative or salt thereof, for moulded capsule, shell, or linker components.

99. (New) The composition according to Claim 98 wherein the dissolution modifying excipient also includes a disintegrant.

100. (New) The composition according to Claim 99 wherein the disintegrant is sodium starch glycollate, croscarmellose sodium, copovidone, or crospovidone (cross-linked polyvinyl pyrrolidone), or a combination or mixture thereof.

101. (New) The composition according to Claim 100 wherein the dissolution modifying excipient also includes a wicking agent.

102. (New) The composition according to Claim 101 wherein the wicking agent is lactose.

103. (New) The composition according to Claim 101 which further comprises a processing aid which is talc.

104. (New) The composition according to Claim 98 wherein the dissolution modifying excipient also includes a wicking agent.

105. (New) The composition according to Claim 104 wherein the wicking agent is lactose.

106. (new) The composition according to Claim 98 wherein the HPC is present in an amount of 10 to about 70% w/w.

107. (new) The composition according to Claim 106 wherein the HPC is present in an amount of 40 to about 70% w/w.

108. (New) The composition according to Claim 107 further comprises a wicking agent which is lactose.

109. (New) The composition according to Claim 107 wherein the lactose is present in an amount of about 0 to 10% w/w.

110. (New) The composition according to Claim 107 wherein the lactose is present in an amount of about 5% w/w.

111. (new) The composition according to Claim 99 wherein the distintegrant is present in an amount of about 10 to about 40% w/w.